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The role of climate factors in geographic variation in body mass and wing length in a passerine bird

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Abstract

Background: Geographic variation in body size is assumed to reflect adaptation to local environmental conditions. Although Bergmann's rule is usually sufficient to explain such variation in homeotherms, some exceptions have been documented. The relationship between altitude, latitude and body size, has been well documented for some vertebrate taxa during the past decades. However, relatively little information is available on the effects of climate variables on body size in birds.

Methods: We collected the data of 267 adult Eurasian Tree Sparrow (*Passer montanus*) specimens sampled at 48 localities in China's mainland, and further investigated the relationships between two response variables, body mass and wing length, as well as a suit of explanatory variables, i.e. altitude, latitude, mean annual temperature (MAT), annual precipitation (PRC), annual sunshine hours (SUN), average annual wind speed (WS), air pressure (AP) and relative humidity (RH).

Results: Our study showed that (1) although the sexes did not differ significantly in body mass, males had longer wings than females; (2) body mass and wing length were positively correlated with altitude but not with latitude; (3) body mass and wing length were negatively correlated with AP and RH, but not significantly correlated with WS. Body mass was positively correlated with SUN and inversely correlated with MAT. Wing length was not correlated with MAT in either sex, but was positively correlated with SUN and negatively correlated with PRC in male sparrows; (4) variation in body mass could be best explained by AP and SUN, whereas variation in wing length could be explained by RH and AP in both sexes. In addition, variation in male sparrows can be explained by SUN, WS and PRC but not in females.

Conclusions: Two different proxies of body size, body mass and wing length, correlated with same geographic factors and different climate factors. These differences may reflect selection for heat conservation in the case of body mass, and for efficient flight in the case of wing length.

Keywords: Body mass, Wing length, Altitude, Latitude, Climate factor, Eurasian Tree Sparrow

Background

Body size is arguably the most important trait affecting the physiology and ecology of animals (Schmidt-Nielsen 1984; Lomolino and Perault 2007; Kingsolver and Huey 2008). Intraspecific, geographic variation in body size is assumed to reflect adaptation to local environmental

conditions (Mayr 1956; Millien et al. 2006; Yom-Tov and Geffen 2011). Bergmann's rule, which postulates that individuals from cooler regions tend to be larger than congeners from warmer regions (McNab 1971), is commonly invoked to explain such variation in both homeotherms (McNab 1971; Blackburn and Ruggiero 2001; Ashton 2002a; Lin et al. 2008; Martinez et al. 2013) and ectotherms (Ashton 2002b; Jin et al. 2006; Pincheira-Donoso et al. 2008; Jaffe et al. 2016). This rule has been corroborated by data on numerous bird and mammal species in the past few decades; however, a few notable

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exceptions have been documented (reviewed by Ashton 2002a and Meiri and Dayan 2003).

Both increasing altitude and latitude are generally negatively correlated with temperature (Blackburn et al. 1999; Potapov 2004; Keller et al. 2013). Previous studies have demonstrated a positive relationship between body size and latitude in both birds and mammals (Ashton 2002a; Meiri and Dayan 2003; Gardner et al. 2009; Olson et al. 2009). However, limited information is available on altitudinal variation in body size (Hamilton 1961; Blackburn and Ruggiero 2001; Chown and Klok 2003; Wilson et al. 2010; Gutiérrez-Pinto et al. 2014). Although latitudinal and altitudinal gradients show similar temperature trends (Ashton and Feldman 2003), some climate factors, such as solar radiation, air pressure (AP) (or oxygen concentration), are more strongly associated with variation in altitude than latitude (Liao et al. 2006; Körner 2007, also see Additional file 1: Table S1). The effects of such climate factors on body size could be as, or more, important than the effects of temperature (Liao et al. 2006; Zhao et al. 2013).

McNab (2010) argued that vertebrates become larger or smaller geographically depending on the abundance, availability and size of resources, and termed this pattern as the “resource rule”. Food availability (i.e. quantity and quality of nutrition, often use net primary productivity as proxy), especially during the growth period, is a crucial predictor determining the final body size, which is impacted directly or indirectly by some abiotic factors such as solar radiation, precipitation, humidity and temperature and others (Rosenzweig 1968; Yom-Tov and Geffen 2011).

It was well known that animals at high altitude must adapt to the stress of lower AP (or oxygen concentration) relative to sea level and still sustain aerobic metabolic processes. Although animals inhabiting higher altitudes generally have higher energy demands for cold surroundings (Snyder 1981; Chappell et al. 1988), limited oxygen availability might decrease digestive efficiency and thus affect negatively their body size. This mechanism has been demonstrated in geographic size variation in lizards and mammals (Jin et al. 2006, 2007; Liao et al. 2006). However, there is little available information on the effect of AP on body size in birds.

Wind speed (WS) has two aspect functions on the body size of an animal. In addition to imposing a thermoregulatory constraint on organisms along with ambient temperature (wind chill effect, Goldstein 1983), strong wind also cause animals, especially flyers, to consume more energy during flight (Bowlin and Wikelski 2008). Therefore, many flying vertebrates and invertebrates modify their wing length or shape to avoid the negative impacts brought by high WS. For example, the elytra and wings of grasshoppers in the Tibetan Plateau have degenerated

or even disappeared (Yin 1984). Wing length of European Storm Petrels (*Hydrobates pelagicus*) tends to increase at high latitudes perhaps in response to local ambient temperature and WS (Jakubas et al. 2014).

Species with large geographic ranges are more likely to exhibit geographic clines in body size (Meiri et al. 2007) because they encounter, and must adapt to, a diverse gradient of environmental conditions throughout their range. This is especially true of sedentary birds (Meiri and Dayan 2003). From this perspective, the Eurasian Tree Sparrow (*P. montanus*), a species resident within a vast geographic range that includes both low and high altitude areas of the Eurasian continent (Fu et al. 1998; Summers-Smith 2009), is an ideal candidate for the study of geographic variation in body size. In this study, we present data on altitudinal and latitudinal variation in body mass and wing length in Chinese populations of the Eurasian Tree Sparrow and use these data to determine whether Bergmann’s rule applies to this species. In addition, we analyze the relationships between body mass, wing length and selected climate factors, including mean annual temperature (MAT), annual precipitation (PRC), annual sunshine hours (SUN), annual average WS, AP and relative humidity (RH) and determine which of these factors are likely to have the greatest effect on body mass and wing length in the Eurasian Tree Sparrow.

Methods

Data collection

This study was based on Eurasian Tree Sparrow specimens housed at the National Zoological Museum of China, Beijing. Body mass, wing length, sex, acquisition time and sites [latitude (°), longitude (°) and altitude (m)] of each individual were extracted from the records of the original collection. In order to ensure the accuracy and reliability of the data, individual birds with worn feathers, or those that were collected during the pre-basic molt stage and breeding stage were excluded from our sample. In the end, we had obtained the body mass and wing length measurements of 267 adult specimens (153 males and 114 females) sampled between 1960 and 1980 at 48 localities in China’s mainland at altitudes ranging from −130 to 4370 m above the sea level and at latitudes ranging from 26°04’N to 52°04’N (Additional file 1: Table S2).

Because body mass and wing length would be expected to be related to climate conditions in the year in which specimens were collected (Yom-Tov et al. 2006), we obtained climate data for the sampling sites in the years in which specimens were collected from the China Meteorological Data Sharing Service System (<http://data.cma.gov.cn/site/index.html>). These climate data included MAT, PRC, SUN, WS, AP and RH (Additional file 1: Table S2). Of these, MAT, PRC, SUN and RH

were significantly correlated with altitude and latitude, whereas AP was only correlated with altitude and WS only with latitude (Additional file 1: Table S1).

Statistical analysis

We conducted one-sample Shapiro–Wilk test and Levene’s test to determine whether data conform to normal distribution and homogeneity of variances. Since body mass and wing length may be affected by gender (Fairbairn et al. 2007), we ran linear mixed-effect models (LME) fitted with the restricted maximum likelihood (REML) method using the *lme* function of *nlme* package in Program R v. 3.2.2 (Pinheiro et al. 2015) to examine the fixed effects of sex (as dummy variable) on body mass and wing length, while accounting for collection site and individual identity as random effects. We further examined the relationship between body mass and wing length for each gender. LME fitted with the REML method was also used to examine the fixed effects of each geographic/climatic factor on body mass/wing length while accounting for collection site and individual identity as random effects. These statistical analyses were performed using Program R v. 3.2.2. The data are presented as mean \pm SE.

We ran a generalized linear model (GLM) using the *glm* function in Program R v. 3.2.2 to model the relationship between body mass or wing length and AP, PRC, SUN, WS, RH and MAT. Because male sparrows had longer wings than females (see “Results” sect.), we analyzed the relationship between wing length and other factors for males and females separately. We used Akaike’s Information Criterion (AIC) to select the best model(s). Multi-parameter models were discarded if a nested model (collinearity among the climate factors), containing a subset of the same parameters, had a better AIC score (Arnold 2010). All possible models within 95% cumulative AIC weight for body mass and wing length were selected and averaged to identify the most important variables using the importance score in the Program R v. 3.2.2 MuMIn package (Kamil 2013).

Results

Although there was no significant difference in body mass between the sexes ($t = -0.444$, $p = 0.675$), male sparrows had longer wings than females ($t = -4.473$, $p < 0.001$; Fig. 1a). Therefore, data on body mass were pooled for analysis whereas wing length data were analyzed separately for each sex (Table 1). Body mass was positively correlated with wing length in each sex (male: $t = 3.856$, $p < 0.001$; female: $t = 4.459$, $p < 0.001$; Fig. 1b). Body mass and wing length in both sexes were positively correlated with altitude, but were not significantly correlated with latitude (Table 1, Fig. 2a, b).

Among the six climate factors examined, body mass and wing length were negatively correlated with AP, and RH, but not significantly with WS in either sex (Table 1; Figs. 3a, b, 4a, b). Body mass was negatively correlated with MAT and positively correlated with SUN (Table 1; Fig. 3c, d). Wing length was not significantly correlated with MAT in either sex, nor with PRC and SUN in females, but was positively correlated with SUN and negatively correlated with PRC in males (Table 1; Fig. 4c, d).

AP, SUN and PRC were the best predictors of body mass, explaining 50% of all variation in this variable (Additional file 1: Table S3). Of these factors, AP and SUN were the most important (Table 2). In males, AP, PRC, RH, SUN and WS were the best predictors of wing length, explaining 50% of all variation in this variable (Additional file 1: Table S3). The relative importance coefficients of these five factors were all greater than 0.6 (Table 2). However, in females, RH, AP and PRC were the best predictors explaining 32% of all variation in this variable (Additional file 1: Table S3). Of these, RH and AP were the most important climate factors (Table 2).

Discussion

Geographic and climatic variation in body mass and wing length

Consistent with Bergmann’s rule, we found that Eurasian Tree Sparrows were heavier at higher altitudes, with a negative correlation between body mass and MAT. This trend has also been reported in a number of sedentary and migratory bird species, such as House Sparrows (*Passer domesticus*; Johnston and Selander 1973), Crested Duck (*Lophonetta specularioides*; Bulgarella et al. 2007), Torrent Duck (*Merganetta armata*; Gutiérrez-Pinto et al. 2014) and some Andean passerine birds (Blackburn and Ruggiero 2001), but not in the Rufous-necked Snowfinch (*Montifringilla ruficollis*; Lu et al. 2009). From a thermoregulatory viewpoint, increased body mass could be an adaptation to colder temperatures (Gardner et al. 2009; Teplitsky and Millien 2014); the reduced surface area to volume ratio of larger-bodied individuals means that they lose proportionately less heat than smaller-bodied birds (Mayr 1956, 1963; Blackburn et al. 1999).

In the present study, we found a positive relationship between wing length and body mass in both sexes, while both surrogates of body size were positively correlated with altitude. In birds, wing length is commonly considered an index of lean body mass (Nolan and Ketterson 1983), so the increased body mass at higher altitudes could be mainly the result of an increase in wing size and flight muscle (Sun et al. 2016), or an increase in the size of metabolic and respiratory organs, i.e. the heart and lungs (Hartman 1955; Carey and Morton 1976; Sun et al. 2016).

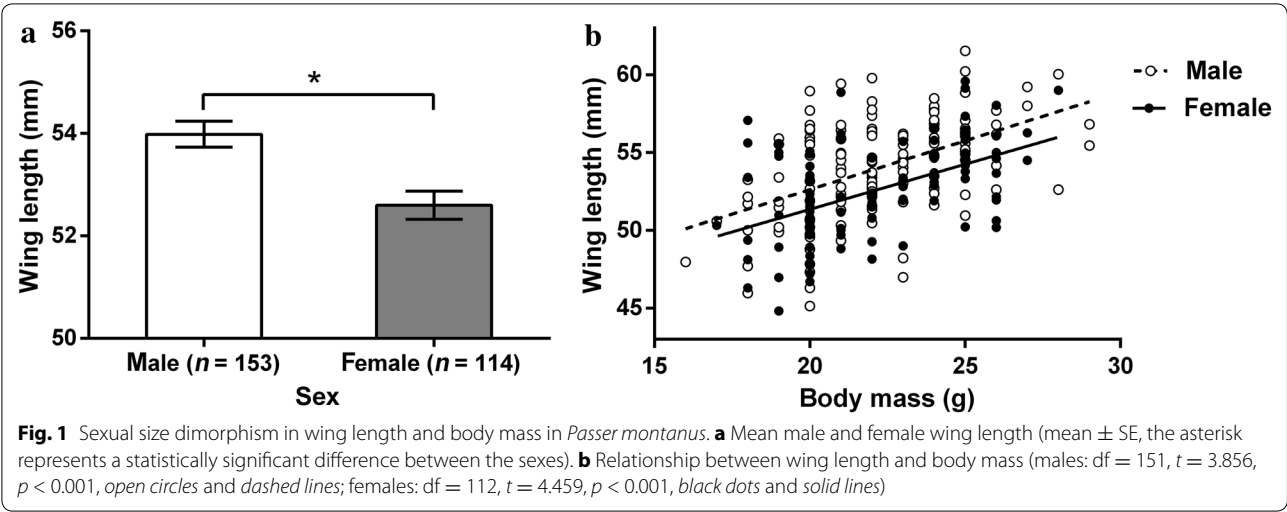
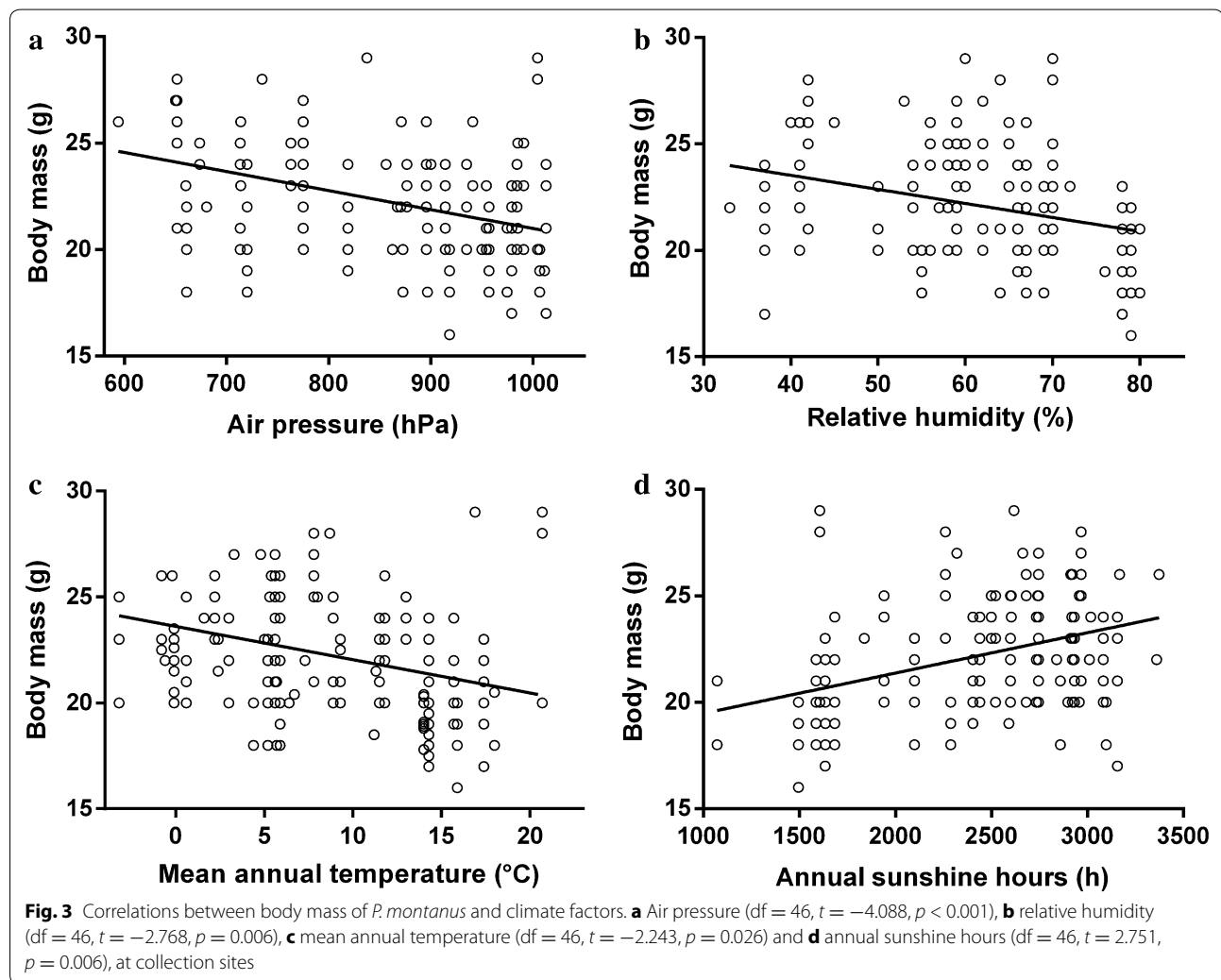
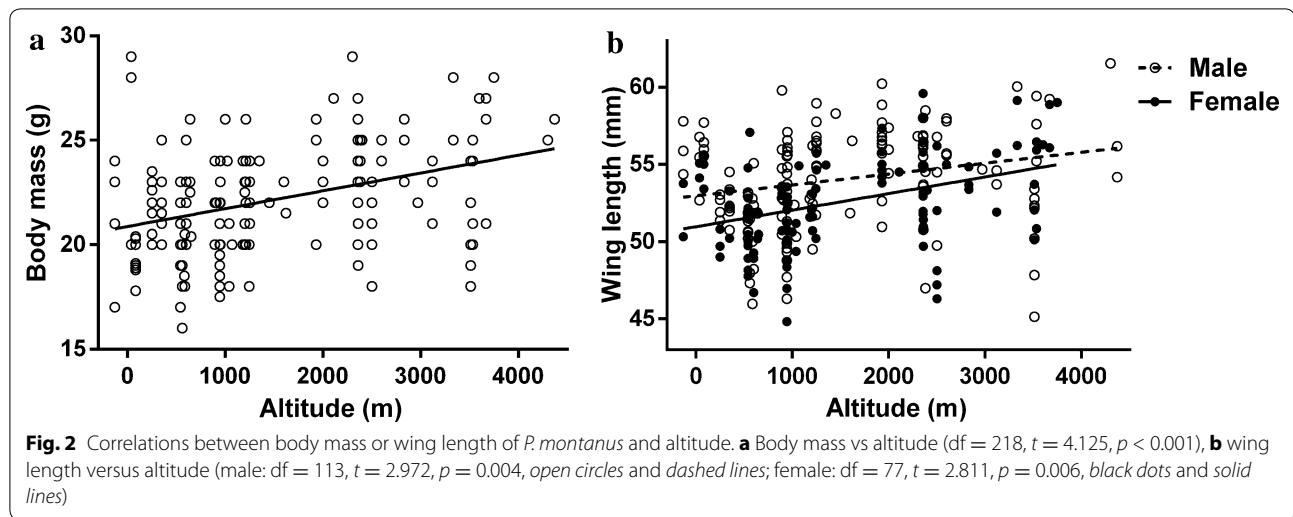


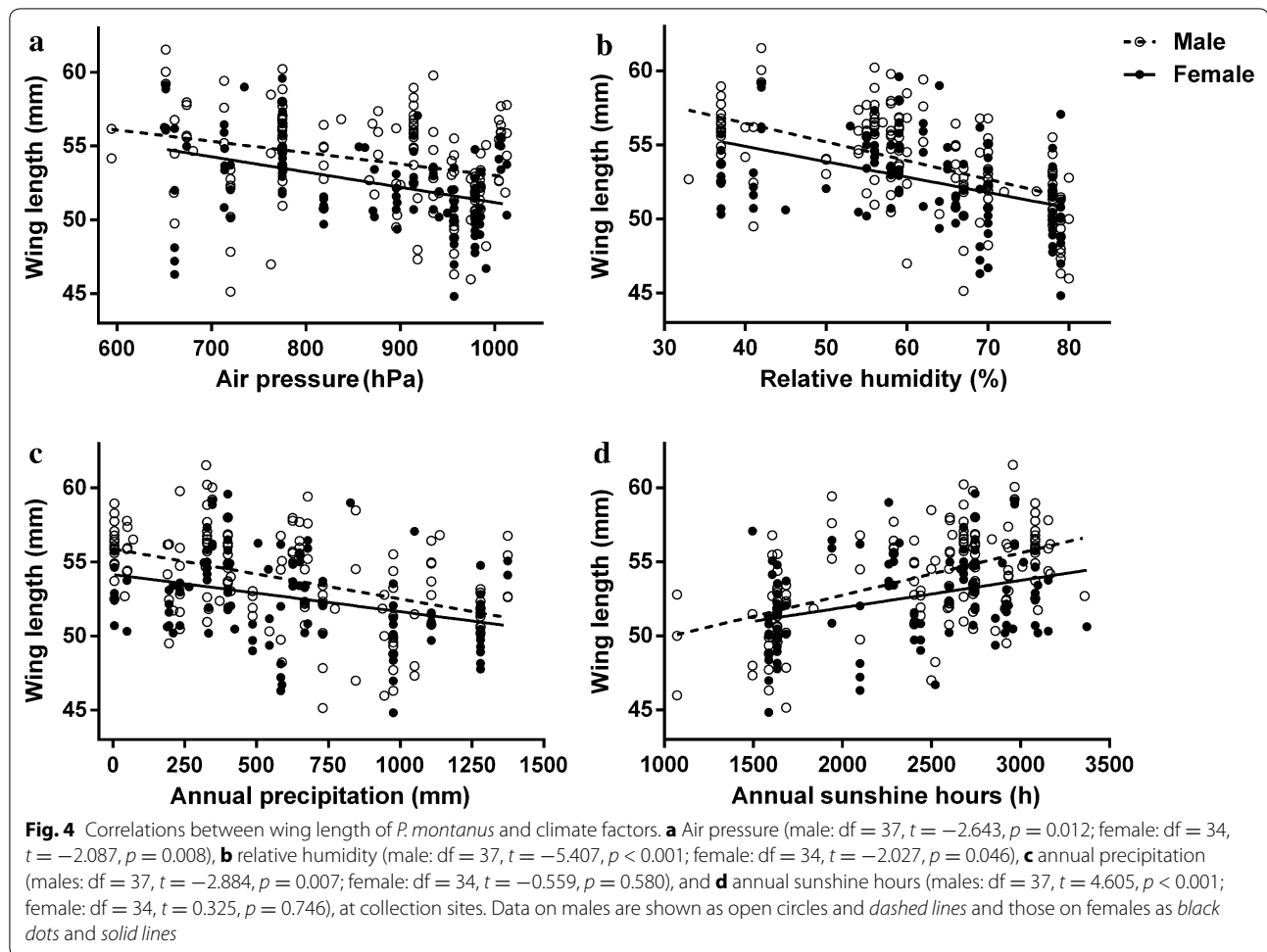
Table 1 Correlations between body mass or wing length of *P. montanus* and geographic and climate factors

Variable	Statistical parameter	Altitude	Latitude	MAT	AP	PRC	SUN	RH	WS
<i>Body mass</i>									
	df	218	218	46	46	46	46	46	46
	Estimate	0.001	−0.026	−0.123	−0.010	−0.001	0.002	−0.070	0.422
	<i>t</i>	4.125	−0.518	−2.243	−4.088	−1.542	2.751	−2.768	1.138
	<i>p</i>	<0.001	0.605	0.026	<0.001	0.130	0.006	0.006	0.261
<i>Wing length</i>									
Male	df	113	113	37	37	37	37	37	37
	Estimate	0.001	−0.052	−0.004	−0.009	−0.003	0.002	−0.129	0.167
	<i>t</i>	2.972	−0.868	−0.060	−2.643	−2.884	4.065	−5.407	0.341
	<i>p</i>	0.004	0.388	0.952	0.012	0.007	<0.001	<0.001	0.735
Female	df	77	77	34	34	34	34	34	34
	Estimate	0.001	−0.123	0.052	−0.009	−0.001	0.000	−0.067	−0.557
	<i>t</i>	2.811	−2.014	0.682	−2.807	−0.559	0.325	−2.027	−1.031
	<i>p</i>	0.006	0.050	0.497	0.008	0.580	0.746	0.046	0.310

Estimate model coefficients, *df* degree of freedom, *MAT* mean annual temperature, *PRC* annual precipitation, *SUN* annual sunshine hours, *WS* annual average wind speed, *AP* air pressure, *RH* relative humidity. Data on the body mass of each sex were pooled but wing length data were analyzed separately for each sex

From the point of view of ambient temperature, animals should exhibit a uniform body size cline along with latitude and altitude (Ashton and Feldman 2003). However, the increasing body size of Eurasian Tree Sparrow was only accompanied with rising altitude, but not with latitude. The different trends in body mass and wing length with increasing altitude or latitude suggest that climate factors other than temperature could have an important effect on body size. For example, variation in AP or oxygen concentration is directly affected by altitude, but rarely by latitude. We found an inverse correlation between AP and both body mass and wing length. As mentioned in the introduction, there is a dilemma that maximum metabolic rates of birds will decrease when they are under low oxygen concentration and increase when they are stressed by low environmental temperatures. The lower wing loadings of longer wings can offset the increased energy requirements incurred by the lower air density at higher altitudes (Swaddle and Lockwood 2003; Altshuler and Dudley 2006). Birds living at higher altitudes typically have a larger heart and lungs in order to provide adequate oxygen to meet metabolic requirements, especially the requirements of flight (Hartman 1955; Carey and Morton 1976; Monge and Leon-Velarde 1991; Scott 2011). The hearts and lungs of Eurasian Tree Sparrows collected at higher altitudes were heavier than those of conspecifics collected at lower altitudes (Sun et al. 2016). This suggests that the positive correlations





between body mass or wing length and altitude may not only reflect adaptations to reduce energy consumption, but also to ensure an adequate oxygen supply.

We found that SUN was positively correlated with both wing length (only for males) and body mass in the Eurasian Tree Sparrow. A sufficient amount of sunshine is conducive to improving food availability by increasing net primary productivity (Churkina and Running 1998; Yom-Tov and Geffen 2011) and raising plant nutrient levels (Potapov 2004). Increased food availability is generally associated with an increase in body size (Yom-Tov and Nix 1986; Aava 2001; Yom-Tov and Gefen 2006, 2011). The Eurasian Tree Sparrow is a typical omnivorous passerine and mainly feeds on grass seeds, fruits and small insects (Fu et al. 1998). Higher sunshine hours per year would not only provide sparrows with more abundant and more nutritious foods, but also extend their available foraging time, thereby making it easier for the birds to accumulate energy and gain body mass.

Unexpectedly, we found that RH was negatively correlated with wing length and body mass and that PRC was negatively correlated with wing length of male sparrows. A similar trend has been reported in the Crested Lark (*Galerida cristata*; Guillaumet et al. 2008). One explanation for this may be the necessity to minimize water loss in cold, dry environments (James 1970; Burnett 1983; Lin et al. 2008). Because the rate of water loss mainly depends on the relative surface area of an organism and its metabolic rate (Chew 1955; Chew and Dammann 1961; Lin et al. 2008), it follows that small-bodied individuals are more vulnerable to acute dehydration than large-bodied ones (Hamilton 1958, 1961; James 1970; Olalla-Tárraga et al. 2009; McKechnie and Wolf 2010).

There was no any correlation between WS and body mass. However, WS is one of most important climate factors for explaining wing length in male but not in female Eurasian Tree Sparrows. Previous studies have reported mixed patterns of the effect of WS on body mass, e.g. positive correlation in Gray Jays (*Perisoreus*

Table 2 Coefficients of averaged models explaining body mass and wing length (male and female) in relation to climate factors

Variable	Factor	Estimate	SE	P value	Lower CI	Upper CI	Relative importance
<i>Body mass</i>							
	(Intercept)	24.181	2.493	<0.001	19.287	29.075	
	AP	−0.008	0.001	<0.001	−0.010	−0.005	1.00
	SUN	0.002	0.001	0.001	0.001	0.003	0.96
	PRC	0.001	0.001	0.453	0.0004	0.003	0.54
<i>Wing length</i>							
Male	(Intercept)	57.864	4.750	<0.001	48.517	67.211	
	SUN	0.002	0.001	0.018	0.001	0.004	0.97
	WS	−0.740	0.428	0.085	−1.537	−0.141	0.88
	RH	−0.099	0.052	0.057	−0.182	−0.051	0.85
	AP	−0.004	0.003	0.149	−0.009	<0.001	0.84
	PRC	0.002	0.001	0.256	0.000	0.005	0.68
	MAT	0.022	0.054	0.688	0.054	0.229	0.15
Female	(Intercept)	66.425	2.666	<0.001	61.158	71.693	
	RH	−0.130	0.060	0.030	−0.248	−0.012	1.00
	AP	−0.006	0.003	0.063	−0.012	−0.003	0.85
	PRC	0.001	0.002	0.502	−0.001	0.007	0.47
	WS	−0.376	0.529	0.479	−1.798	0.118	0.45
	MAT	−0.035	0.089	0.691	−0.384	−0.084	0.15

Estimate average model coefficients, SE unconditional standard errors, Lower CI and Upper CI the 95% confidence limits, AP air pressure, PRC annual precipitation, SUN annual sunshine hours, RH relative humidity, WS annual average wind speed, MAT mean annual temperature

More information for top-ranked models are shown in Additional file 1: Table S3

canadensis; Waite 1992), negative correlation in European Storm Petrels (*Hydrobates pelagicus*; Jakubas et al. 2014), or no correlation in the Dovekie (*Alle alle*; Wojczulanis-Jakubas et al. 2011). In general, relatively longer wings can increase flight efficiency (Lockwood et al. 1998); many species, especially in migratory or insectivorous birds, are characterized by long wings for gliding and soaring in strong air currents (Landmann and Winding 1995; Lockwood et al. 1998; Nowakowski 2000). Given that little information is available on the effects of WS and flight ability for this sedentary passerine bird, further investigation of such interspecific and gender differences in flight efficiency are needed to clarify the inconsistent results.

Among the six climate factors, only AP and SUN were important with respect to variation in body mass. RH and AP were the most important with respect to variation in wing length in both male and female sparrows. In addition, SUN, WS and PRC were the most important in males but not in females. Therefore, altitudinal and latitudinal variation in body mass and wing length in the Eurasian Tree Sparrow appears to be affected by a combination of climate factors and sexual selection. Unexpectedly, although there was a significant correlation between body mass and MAT, the latter was not a

reliable predictor of body mass in our statistical model. Since we were not able to obtain a satisfactory answer, we deem it necessary to conduct further research on this aspect.

Sexual size dimorphism in body mass and wing length

The Eurasian Tree Sparrow is generally considered sexually monomorphic. However, although we found no significant difference in body mass between the sexes, males had longer wings than females. These results are consistent with those obtained for other populations or other subspecies of the Eurasian Tree Sparrow (St. Louis and Barlow 1991; Mónus et al. 2011; Sun et al. 2016). Although sexual displays in birds require much energy (Gil and Gahr 2002; Ward and Slater 2005), male Eurasian Tree Sparrows can behave in the same intensity during autumnal sexual recrudescence as they commonly do at pre-breeding (García-Navas et al. 2008; Pinowski et al. 2009). That is, as in the spring, males sing and display in the fall, compete for territories, attract females, build nests and copulate with females (García-Navas et al. 2008; Pinowski et al. 2009). The longer wings in males are thought to reduce the flight costs incurred in sexual display (Møller 1991; Hedenstrom and Møller 1992). Therefore, sexual size dimorphism in wing length in the Eurasian Tree Sparrow

may reflect male-specific reproductive behavior and in spite of it, this may also covary with a number of other morphological traits (see Table 1 in Mönus et al. 2011). Furthermore, female birds are more susceptible to predators because they incubate or brood the nest, although the short wings of some female passerines may enhance their maneuverability and help them escape predation (Swaddle and Lockwood 2003; Bomberger and Brown 2011).

Conclusions

Geographic variation in body mass and wing length in the Eurasian Tree Sparrow is generally consistent with Bergmann's rule. Body mass and wing length is positively correlated with altitude, but not with latitude, suggesting that the body size of sparrows is more affected by other climate factors than by MAT. Most variation in body mass can be explained by AP and SUN, whereas significant variation in wing length can be best explained by RH and AP in both males and females. In addition, variation in male sparrows can be explained by SUN, WS and PRC but not in females. Two proxies of body size, i.e. body mass and wing length, displayed different strengths and polarities of correlation with the same geographic and climate factors. These differences may reflect competing selection pressures for heat conservation, flight efficiency and sexual selection.

Additional file

Additional file 1: Table S1. Partial correlations between the altitude and latitude of Eurasian Tree Sparrow (*Passer montanus*) collection sites and selected climatic factors. **Table S2.** Sampling information of Eurasian Tree Sparrows and climate data of their collection sites. **Table S3.** Top-ranked cumulative link models of the relationship between body mass or wing length of the Eurasian Tree Sparrow and potential explanatory factors.

Authors' contributions

YFS, DML and YFW conceived the research project, ML and GS collected the data, YFS analyzed the data and YFS and DML led the writing with help from FML. All authors have read and approved the manuscript.

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Competing interests

The authors declare that they have no competing interests.

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